201-16303B

I. General Information

CAS Number:

C.I. Pigment Violet 19, (CAS No. 1047-16-1)

RECEIVED OPPT C810

Name:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro

06 JUN 28 AM 11:59

CAS Number:

C.I. Pigment Violet 122, (CAS No. 980-26-7)

Name:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro-2,9-dimethyl

CAS Number:

(CAS NO. 5862-38-4)

Name:

Dihydro Quinacridone

II. Physical-Chemical Data

A1. Melting Point

Test Substance

Test substance:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro

Remarks:

Method

Method:

Measured

Remarks:

Results

Melting point value:

>400 °C

Remarks:

References

Other

Anliker R. and Moser P., The Limits of Bioaccumulation of Organic Pigments in Fish: Their Relation to the Partition Coefficient and the Solubility in Water and Octanol, Ecotox. And Envir. Saf. 13, Pp. 43-52 (1987) Data is consistent with melting points for the class of pigments and other available measurements,

A2. Melting Point

Test Substance

Test substance: Quino(2,3-b)acridine-7,14-dione,5,12-dihydro-2,9-dimethyl

Remarks:

Method

Method: Adapted Joback Method

Remarks:

Results

References

Melting point value: 349 °C

Remarks:

EPIWIN v 3.10, Syracuse Research Corporation, Syracuse, New York

Other Data is consistent with melting points for the class of pigments and other

available measurements.

A2. Melting Point

Test Substance

Test substance: Dihydro Quinacridone

Remarks:

Method

Method: Esitimate, Adapted Joback method

Remarks:

Results

Melting point value: 349.84 °C

Remarks:

References EPIWIN v 3.10, Syracuse Research Corporation, Syracuse, New York

Other Data is consistent with melting points for the class of pigments and other

available measurements.

В.	Boiling Point Test Substance Test substance: Remarks:	SOLID N/A
	Method Method: Remarks:	
	Results Boiling point value: Remarks:	
	References	
	Other	
C1. Vapor Pressure Test Substance Test substance:		Quino(2,3-b)acridine-7,14-dione,5,12-dihydro
	Remarks:	
	Method Method: Remarks:	Estimation Modified Grain method
	Results Vapor pressure val Temperature:	ue: 1.13 E-010 mmHg
	Remarks:	
	References Other	MPBPWIN v1.40 in EPIWIN v 3.10, Syracuse Research Corporation Syracuse, New York
	apor Pressure	O to (0.0.1) with 7.14 than 5.10 dibudes 2.0 dimethal
Test Substance Remarks:		Quino(2,3-b)acridine-7,14-dione,5,12-dihydro-2,9-dimethyl
Resul		2.14 E-011 mm Hg

Remarks:

Method

Method:

Estimation

Remark:

Modified Grain method

References

York

MPBPWIN v 1.40 in EPIWIN v 3.10, Syracuse Research Corporation, Syracuse, New

D. Partition Coefficient

Test Substance

Test substance: Quino(2,3-b)acridine-7,14-dione,5,12-dihydro

Remarks:

Method

Method: Remarks: Octanol Solubility Determination GLP \1996\ Guideline 40 CFR 796

Results

Solubility:

.808 mg/L at 20 °C

Remarks:

References

Corning Hazleton, CHW 6623-105, 1996, Log Kow partition coefficient cannot be determined for this compound, solubility in water and octanol are

too low to produce a meaningful value.

Other

E. Water Solubility

Test Substance

Test substance:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro

Remarks:

Method

Method:

Estimated

Remarks:

Results

Value:

<.808 mg/L

Temperature:

20 °C

Description:

Remarks:

Extremely Low Solubility

References

Corning Hazleton, CHW 6623-105, 1996, Log Kow partition coefficient

cannot be determined for this compound, solubility in water and octanol are too

Other low to produce a meaningful value.

III. Environmental Fate Endpoints

A. Photodegradation

Test Substance

Test substance:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro

Remarks:

Method

Method: Test type:

Remarks:

Estimate

Water\sunlight

Results

Temperature: Degradation Rate

Half-life

.642 hours

Ozone reaction:

Remarks:

Conclusions

[Estimate only applies to minute soluble fraction]

References

AOPWIN v 1.91, Syracuse Research Corporation, Syracuse, New York

A2. Photodegradation

Test substance:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro-2,9-dimethyl

Remarks:

Method

Method:

Estimate

Test type: Water\sunlight

Remarks:

Results

Temperature:

Hydroxyl radicals reaction OH Rate constant:

Half-life

.641 hours

Ozone reaction:

Remarks:

Conclusions

[Estimate only applies to minute soluble fraction]

References

AOPWIN v 1.91, Syracuse Research Corporation, Syracuse, New York

B. Stability in Water

Test Substance

Test substance:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro

Remarks:

Method

Method: Test type: GLP: Remarks:

Results

Half-life:
Percent hydrolyzed in
5 days (120 hs)
at 50 °C:
Remarks:

Conclusions

Data Quality

Remarks:

References

Other

Due to extremely low solubility, hydrolosis in water for quinacridone pigments cannot be estimated or measured accurately at this time. See HYDROWIN v 1.67 Syracuse Research Corporation, Syracuse, New York

C. Biodegradation

Test Substance

Test substance:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro

Remarks:

Method

Method:

Estimation

Test type: GLP: Year: Remarks:

Results

Results:

noreadily biodegradable

Remarks:

Conclusions

Results apply to all three quinacridone pigments.

Data Quality

Remarks:

References

EPI Suite HYDROWIN v 4.02 Syracuse Research Corporation, Syracuse, New

York, Anliker R., and Clarke, E.A. Ecology and Toxicology of Synthetic

Organic Pigments, Chemosphere, Vol. 9, pp. 595-609 (1980)

D. Transport between Environmental Compartments (Fugacity)

Test Substance

Test substance:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro

Remarks:

Method

Test type:

Estimation

Model used:

Level III Fugacity Model; EPIWIN:EQC from Syracuse Research

Corporation

Remarks:

Results

Model data and results:

Distribution (%)

 Air
 5.15 E-007

 Water
 37.1

 Soil
 62.8

 Sediment
 .0897

Remarks:

Since no experimental values were available the physical chemical values

utilized in this model were default parameters from within EPIWIN.

References

Conclusions

Meylan, W. (1993). User's Guide for the Estimation Programs Interface (EPI), Version 3.10, Syracuse Research Corporation, Syracuse, New York 13210. The Level III model incorporated into EPIWIN is a Syracuse Research Corporation adaptation of the methodology described by Mackay *et*

Other

al. 1996; Environ. Toxicol. Chem. 15(9), 1618-1626 and 1627-1637.

D2. Transport between Environmental Compartments (Fugacity)Test Substance

Test substance:

Remarks:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro-2, 9 dimethyl Estimation Level III Fugacity Model; EPIWIN: EQC from Syracuse Research Corporation

15

Distribution (%) Air

1.56 E-006

Soil

Water 84.8

Sediment

Method Test type:

Model used:

Remarks:

Results Model data and results:

Estimated distribution and media concentration

(levels II/III):

Remarks:

Conclusions

References

Other

.122Since no experimental values were available the physical chemical values utilized in this model were default parameters from within EPIWIN.Meylan, W. (1993). User's Guide for the Estimation Programs Interface (EPI), Version 3.10, Syracuse Research Corporation, Syracuse, New York 13210. The Level III model incorporated into EPIWIN is a Syracuse Research Corporation adaptation of the methodology described by Mackay et al. 1996; Environ. Toxicol. Chem. 15(9), 1618-1626 and 1627-1637.

D3. Transport between **Environmental Compartments** (Fugacity)Test Substance Test

substance:

Remarks:

Method Test type:

Model used: Remarks:

Results Model data and results: Estimated distribution and media concentration (levels II/III):

Remarks:

Conclusions

References

Other

Quino(2,3-b)acridine-7,14-dione,5, 6, 12, 13-tetrahydroEstimationLevel III Fugacity Model; EPIWIN:EQC from Syracuse Research Corporation

Distribution (%)

Air

1.04 E-009

Water 86.1

10.8

Soil

Sediment

3.08Since no experimental values were available the physical chemical values utilized in this model were default parameters from within EPIWIN.Meylan, W. (1993). User's Guide for the Estimation Programs Interface (EPI), Version 3.10, Syracuse Research Corporation, Syracuse, New York 13210. The Level III model incorporated into EPIWIN is a Syracuse Research Corporation adaptation of the methodology described by Mackay et al. 1996; Environ. Toxicol. Chem. 15(9), 1618-1626 and 1627-1637.

IV. Ecotoxicity

A. Acute Toxicity to Fish

Test Substance Quino(2,3-b)acridine-7,14-dione,5,12-dihydro

Test substance: Remarks:

Method

Method: Estimation

Test type: GLP: Year:

Species/strain: Fish

Analytical monitoring: Exposure period: Remarks:

Results
Nominal concentration:

Measured concentration:

Endpoint value:

Biological observations:

Statistical methods:

Remarks:

Conclusions Due to its insolubility, the material is not anticipated to be toxic in the water at

LC50 96 Hours 885 mg/L, 14 Day LC 50 1454 mg/L

saturation.

Data Quality

Reliability: Remarks:

References

EPI Suite ECOSAR v .099 Syracuse Research Corporation, Syracuse, New

York, Anliker R. and Moser P., The Limits of Bioaccumulation of Organic

Other Pigments in Fish: Their Relation to the Partition Coefficient and the Solubility in

Water and Octanol, Ecotox. And Envir. Saf. 13, Pp. 43-52 (1987)

A2. Acute Toxicity to Fish

Test Substance

Test substance:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro-2,9-dimethyl

Remarks:

Method

Estimation

Method: Test type: GLP:

Year:

Fish

Species/strain:

Analytical monitoring: Exposure period:

Remarks:

Results

Nominal concentration:

Measured concentration:

LC50 96 Hours 91 mg/L, 14 Day LC 50, 178.04 mg/L

Endpoint value:

Biological observations:

Statistical methods:

Remarks:

Conclusions Due to its insolubility, the material is not anticipated to be toxic in the water

at saturation.

Data Quality

Reliability: Remarks:

References EPI Suite ECOSAR v .099 Syracuse Research Corporation, Syracuse, New

York, Anliker R. and Moser P., The Limits of Bioaccumulation of Organic

Pigments in Fish: Their Relation to the Partition Coefficient and the Solubility

Other in Water and Octanol, Ecotox. And Envir. Saf. 13, Pp. 43-52 (1987)

Б,	Test substance:	
	Remarks:	Quino(2,3-b)acridine-7,14-dione,5,12-dihydro-2,9-dimethyl
Methoc Ex _l	Method: Test type: GLP: Year: Species/strain: Analytical monitoring: posure period: Remarks:	OECD 211 Daphnia Magna reproduction Yes Daphnia Magna
Results	Nominal concentration: Measured concentration: Endpoint value: Reproduction Biological observations: Statistical methods: Remarks:	No differences in the onset of brood production observed in the concentration group in comparison to the control. The reproduction rate in the concentration group showed no statistically significant changes in comparison to the control.
Conclu	nsions	
Data Ç	Quality Reliability: Remarks:	
Refere	ences	Reliable without restriction
Other		Company sponsored data

C. Toxicity to Aquatic Plants

Test Substance

Test substance:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro

Remarks:

Method

Method:

Estimation

Test type: GLP:

Year:

Species/strain:

Algae

Endpoint basis: Exposure period: Analytical procedures:

Nominal concentration:

Remarks:

Results

The conduction of an algae test with C.I. Pigment Violet 19, Red 122 or dyhydroquinacridone is problematic as the substance leads to a strong

coloring of the test solution and therefore to a reduction of light intensity. Therefore, the assessment is made on the basis of computer model estimation.

Measured concentration:

Endpoint value:

NOEC: Biological

96 hour EC-50, 548.6

observations:

Was control response

satisfactory:

Statistical Methods:

Remarks:

Conclusions

Data Quality

Reliability: Remarks:

References

reliable with restriction

Other

EPI Suite ECOSAR v .099 Syracuse Research Corporation, Syracuse, New

York,

V. Toxicological Data

Acute Toxicity A.

Test Substance

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro Test substance:

Purity was unknown

Remarks:

Method

Acute lethality; Other Method:

LD₅₀ estimate Test type: No (Pre-GLP) GLP:

1957 Year:

Species/strain: Male albino Rats Route of exposure: Oral gavage

1000, 3400, 5000, 7500 mg/kg Dose levels: Remarks:

Results

 $LD_{50} = >7,500 \text{ mg/kg}.$ Value:

Deaths at each dose:

Remarks: All rats survived, Clinically, the rats showed only mild discomfort at the

higher levels. The material appeared to be excreted in the feces.

Conclusions

Material would be considered as not toxic.

Data Quality

Reliability: Reliable with restrictions

Remarks: The study was conducted quite some time ago and hence many study details

are missing from the report and not available. However, basic data are given

and results are consistent with other data for pigments of this type.

Haskwll Laboratory, Medical Research project, No., MR-166, See also, Mone References

> J.G. 1968, Federation Series on Coating Technology, Unit 9 Organic Pigments, Federation of Societies for Paint Technology, Philadelphia, PA

Other 19107. **Acute toxicity**

Test substance:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro-2,9-dimethyl

Remarks:

Purity was unknown

Method

Method:

Acute lethality; Other

Test type:

LD₅₀ estimate

GLP:

No (Pre-GLP)

Year:

1968

Species/strain: Route of exposure: Rat and mouse Oral gavage

Dose levels:

Unknown

Remarks:

Results

 $LD_{50} = >5,000 \text{ mg/kg}.$

Deaths at each dose:

Remarks:

Value:

Conclusions

Material would be considered as not toxic.

Data Quality

Reliability:

Reliable with restrictions

Remarks:

References

Mone J.G. 1968, Federation Series on Coating Technology, Unit 9 Organic

Pigments, Federation of Societies for Paint Technology, Philadelphia, PA

19107.

A. Acute Toxicity

Test Substance

Test substance: Quino(2,3-b)acridine-7,14-dione,5,12-dihydro

Purity was unknown

Remarks:

Method

Method: Acute lethality; Other

 $\begin{array}{ll} \text{Test type:} & \text{LC}_{50} \text{ estimate} \\ \text{GLP:} & \text{No (Pre-GLP)} \end{array}$

Year: 1983

Species/strain: Male CRL:CD® Rats

Route of exposure: Inhalation

Dose levels: 1.5, 1.6, 2.4, 2.6 and 3.1 mg/l

Remarks:

Results

Value: $LC_{50} = >3.1 \text{ mg/L}$

Deaths at each dose:

Remarks: All rats survived, Groups of 6 rats were used at each dose up to 3.1 mg/L.

Other than transint weight losses there were no significant clinical signs of

toxicity observed.

Conclusions

Material would be considered as not toxic.

Data Quality

Reliability: Reliable with restrictions

Remarks: The study is well documented and followed accepted protocols.

References Haskell Laboratory, Medical Research Report Number 746-82, Project, No.,

MR-4368-001,

Repeated Dose Toxicity Test

Substance

Test substance:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro

Remarks:

Method

Method:

Repeated subchronic dose

Test type:

GLP: Year: NA 1982

Species/strain:

Fisher 344 Rats

Route of exposure:

Gavage

Duration of test:

33 days

Exposure levels:

Rats 0. 1.0%, 5.0 %, 10.0% in the diet

Sex:

Exposure period:

33 days

Post-exposure Observation period:

Remarks:

Results

NOAEL (NOEL):

Up to 10 % of the diet

After repeated oral administration for 33 days in rats, pigment Violet 19 showed no signs of toxicity. None of the study animals died on test. Clinically, high dose (10%) animals demonstrated significant body weight

gain compared to controls, which appeared to be associated with corresponding increase in food intake. It appeared that these animals tried to compensate by overeating for the decrease in nutritional intake in the 10% pigment diet. These animals, and to a lesser extent the 5% and 1% dose level animals, also had purple tinged fur, apparently as a result from coming in contact with the color pigment in feed hoppers. No other clinical sign were seen in the animals. Clinical pathology, ophthalmology, cytogenetic analysis, organ weights, and gross and tissue morphology examinations failed to detect the toxicity associated with Pigment Violet 19. (A very slight but statistically significant increase in methemoglobin levels was seen for the high dose female rats at week 2, but in neither sex at week 4. Not considered related to Pigment Violet 19 treatment.) In general, under the conditions of the study. toxicity was not observed following the administration of up to 10% Pigment

Violet 19 in the diet of Fisher 344 rats for 33 days.

Conclusions

Test substance is not toxic

Data Quality

Reliability:

Reliable without restriction

Remarks:

References:

Microbiological Associates, September, 1988 Study for CTFA,

CTFA 86-MAI-A; MAG1003-T03022, Subchronic Oral Toxicity In Rats.

Repeated Dose Toxicity Test

Substance

Test substance:

Ouino(2,3-b)acridine-7,14-dione,5,12-dihydro

Remarks:

Method

Method:

Absorption/ Distribution/Excretion

Test type:

GLP:

NA 1991

Year: Species/strain:

Fisher 344 Rats

Route of exposure:

Gavage

Duration of test:

72 Hours

Exposure levels:

3.22 mg/kg and 33.68 uCi/kg Males, 5.44mg/kg 56.81 uCi/kg Females

Sex:

Exposure period:

single dose

Post-exposure

72 hour follow up

Observation period:

Remarks:

Results

NOAEL (NOEL):

N/A

The test article was administered as a suspension in aqueous 1%

carboxymethyl cellulose at a concentration of .3905 mg QV19 and the same amount was administered to each rat Urine and feces were collected from

each rat at 2,8,24,48 and 72 hours after dosing; cage washes and

gastrointestinal tract of each rat were removed after euthanasia at 72 hour post-dose. Recovery of administered radioactive dose was virtually complete.91.9+ or - 6.9 % of dose males; 100.5+ or 8.7% of dose females. There were no gender related differences in the route of excretion. More than 90 % of the recovered radioactivity was eliminated in the feces and cage washes, which appeared to contain residual fecal matter. At 72 hours virtually all radioactivity had been eliminated by the rats. The urine from both groups of rats contained very low amounts of radioactivity.0089% of

dose males; .0020% of dose females.

Conclusions

Radioactivity from a single oral dose of Pigment Violet 19 given to male and female rats

was eliminated almost completely in the feces.

Data Quality

Reliability: Remarks:

Reliable without restriction

References:

Bio-Research 1991, Study done for CTFA,

Repeated Dose Toxicity Test

Substance

Test substance:

Ouino(2,3-b)acridine-7,14-dione,5,12-dihydro

Remarks:

Method

Method:

Whole Body Radiography

Test type:

GLP: Year: NA 1991

Species/strain:

Fisher 344 Rats

Route of exposure:

Gavage

Duration of test:

48 Hours

Exposure levels:

??

Sex:

Exposure period:

Post-exposure

single dose 48 hour follow up

Observation period:

Remarks:

s:

Results
NOAEL (NOEL):

N/A

Groups of male and female Fisher 344 rats were administered orally by gavage pigment violet 19 and radioactive trace material. And the tissue distribution of radioactivity determined by whole body autoradiography at selected times up to 48 hours after dosing. The autoradiogram showed that radioactivity was localized only in the gastrointestinal tract of both male and female rats. No radioactivity was detected in other organs and tissues of the animals. The highest concentrations of radioactivity were found at 2 hours post dosing. Most of the radioactivity was eliminated from the rats at 24

hours and it was virtually undetected at 18 hours post-dose.

Conclusions

Whole body autoradiography indicated that virtually no radioactivity was detected in tissues, supporting the previous finding that, radioactivity from a single oral dose of Pigment Violet 19 given to male and female rats was eliminated almost completely in the

feces.

Data Quality

Reliability: Remarks: Reliable without restriction

References:

Bio-Research 1991, Study done for CTFA,

C. Genetic Toxicity - Mutation

Test Substance

Test substances:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro

Remarks:

Method

Method:

In Vitro Mutagenicity\

Test type: GLP:

Ames ?? 1975

Year:

Salmonella typhimurium

Species/strain: Metabolic activation:

Yes

Concentration tested:

103,

Concentration

100 ug per plate

Remarks:

Results

Result:

Negative

Cytotoxic

concentration:

Precipitation

concentration:

Genotoxic effects

With activation:

Negative Negative

Without activation: Statistical methods:

Remarks:

Conclusions

Data Quality

Reliability:

Reliable without restrictions

Remarks:

Six crystal forms of Violet 19 were tested, No mutagenic response was seen

with any of the pigments tested.

References

Salmonella/ Mammalian- microsome plats incorporation mutagenicity/Haskell Laboratory Report No. 558-75, See also CTFA Report, Quinacridone Violet

19

C. Genetic Toxicity - Mutation

Test substance:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro-2,9-dimethyl

Remarks:

Method

Method: Test type: **OECD 471** Ames

GLP: Year:

22 2000

Species/strain:

Salmonella typhimurium

Metabolic activation:

With and without

Concentration tested:

??5000 ug/plate with and without activation

Remarks:

Results

Result:

Negative in all bacterial strains with and without activation

Cytotoxic concentration:

Precipitation concentration:

Genotoxic effects

With activation: Without activation: Negative Negative

Statistical methods:

Remarks:

Conclusions

Data Quality

Reliability:

Reliable without restriction

Remarks:

References

Notox Project No. 289845

Genetic Toxicity - Chromosomal Aberrations D.

Test Substance

Test substance:

Quino(2,3-b)acridine-7,14-dione,5,12-dihydro

Remarks:

Method

Method:

OECD 473??

Test type:

Cytogenetics Assay

GLP:

Year:

2001??

Species/strain:

Exposure period:

Remarks:

Mouse Lymphoma L5178Y Cells

Results

Result:

Negative

Genotoxic effects:

Negative ?????ug/plate

Concentration tested

Statistical methods: Remarks:

Conclusions

Not mutagenic

Data Quality

Reliability:

Reliable without restriction

Remarks:

References

CTFA Micronucleus in vivo and mouse lymphoma cell mutation

underway January, 2000

E. Developmental Toxicity

Test Substance

See subchronic toxicity and absorption studies above.

Test substance:

Remarks:

Method

Method:

GLP:

Year:

Species/strain:

Sex:

Route of exposure:

Exposure levels:

Actual doses received:

Exposure period:

Duration of test:

Remarks:

Results

Maternal toxicity

NOEL:

NOEL for

teratogenicity:

NOEL for fetotoxicity:

Parental toxic

responses:

Fetal toxic responses

dose:

Statistical Methods:

Remarks:

Conclusions

Since available radiographic studies establish consistently no significant uptake or absorption from this substance, no further reproduction or developmental studies are planned.

Data Quality

Reliability: Remarks:

References

F. Toxicity to Reproduction

Test Substance

Test substance:

Remarks:

Method

Method:

GLP:

Year:

Species/strain:Sex:

Route of exposure:

Exposure levels:

Exposure period:

Duration of test:

Remarks:

Results

Maternal toxicity NOEL:

Parental toxic responses:

Fetal toxic responses dose:

Statistical Methods:

Remarks:

Conclusions

Data Quality

Reliability:

Remarks:

References

				• .
Аc	ute	to	K I C	IIV

Test substance:

(1) Quino(2,3-b)acridine-7,14-dione,5,12-dihydro and (2) Quino(2,3-b)acridine-7,14-

dione,5,12-dihydro-2,9-dimethyl

Remarks:

Method

Method:

Skin irritation to the rabbit

Test type:

Skin irritation

GLP:

unknown

Year:

(1)1992 (2) 1982

Species/strain:

Route of exposure:

Dose levels: Remarks:

rabbitt

Results

Value:

negative

Deaths at each dose:

Remarks:

Conclusions

Data Quality

Reliability:

unassignable

Remarks:

References

(1) Dupont Haskell Report HLO 584-82

(2) MB Research Labs Project No. MB 92-1750CD

Other

\\Sbs2003\users\KatieSherman\Test Plans\CIPigment Violet 19 Red122 Dihydro Quin 05 23 06.rtf